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**Potassic late orogenic Stephanian volcanism in the South West French Massif central (Decazeville, Figeac, Lacapelle-Marival basins): an example for mantle metasomatism along strike-slip faults?**

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**Mots clefs:** *magmatisme tardi-orogénique, shoshonitique, métasomatisme, Sillon Houiller, Varisque*

**Abstract**

In the Southwestern part of the French Massif central (Decazeville basin, at the Sillon Houiller fault termination; Figeac and Lacapelle-Marival basins along the Argentat fault), Stephanian volcanism exhibits shoshonitic affinities. Their chondrite-normalized Rare Earth Element (REE) patterns are enriched in light REE, but almost flat for heavy REE, with marked negative Eu anomalies. Primitive mantle-normalized element spectra show negative Nb, Ta, P, Sm, Ti, and positive Th, U, Pb anomalies, respectively.  $\epsilon_{Nd}$  values are negative and homogeneous (-6 to -4). This volcanism shares the same geochemical patterns as the late-

orogenic Stephanian-Permian magmatism from the southern part of the Variscan belt (Pyrénées, Alps, Sardinia). We explain these geochemical characteristics as resulting from the partial melting of a metasomatised mantle. We propose a new mechanism to explain this melting process: horizontal displacement along the main late-orogenic strike-slip faults might bring into contact a hydrated lower crust with the lithospheric mantle. Mantle metasomatism within the strike-slip fault zone may then induce partial melting.

### **Résumé**

*Le volcanisme Stéphalien du Sud Ouest du Massif central français, observé dans les bassins de Decazeville à l'extrémité du Sillon Houiller, de Figeac et Lacapelle-Marival le long de la faille d'Argentat, présente des caractéristiques géochimiques similaires. Il s'agit d'un magmatisme d'affinité shoshonitique, présentant des spectres de terres rares normalisés aux chondrites enrichis en terres rares légères, mais presque plat pour les terres rares lourdes, avec une anomalie négative en Eu marquée. Les spectres élémentaires normalisés au manteau primitif montrent des anomalies négatives en Nb, Ta, P, Sm et Ti, et des anomalies positives en Th, U et Pb. Les valeurs d' $\epsilon_{Nd}$  sont négatives et également très homogènes (-6 à -4). Ce volcanisme présente les mêmes caractéristiques que le volcanisme tardi-orogénique stéphano-permien du Sud de la chaîne varisque (Pyrénées, Alpes, Sardaigne). Ces caractéristiques sont interprétées comme résultant de la fusion partielle d'une source mantellique métasomatisée. Il est proposé un nouveau mécanisme provoquant cette fusion, par l'intermédiaire du déplacement horizontal des failles décrochantes tardi-orogéniques majeures, qui pourrait mettre en contact une croûte inférieure hydratée et le manteau lithosphérique. La métasomatose du manteau dans la zone de faille décrochante provoquerait sa fusion.*

## **Introduction**

Late orogenic times are key periods in mountain belt evolution, when the continental lithosphere acquires a new structure that will control further evolution, including its reworking in new orogens. The continental crust is particularly modified by late orogenic magmatism and tectonism. Bimodal magmatism is widespread, and results primarily from partial melting of the crust and underlying mantle [Bonin, 2004]. The magmas evolving from mantle melts are commonly potassium-rich [High Potassium Calc Alkaline: HKCA, e.g. Liégeois *et al.*, 1998], but the mechanisms associated with this enrichment are still a matter of discussion [Gibson *et al.*, 1993; Liégeois *et al.*, 1998; Miller *et al.*, 1999; Pecerrillo and Martinotti, 2006]. This magmatism is coeval with a change in the tectonic regime, from regional compression to strike-slip, then extensional regime, that precedes the tectonic quiescence [e.g. Blès *et al.*, 1989]. The strike-slip regime generates numerous vertical faults that cross-cut the orogenic structures.

At the end of the Variscan orogeny, Westphalian synkinematic plutonism accommodated extension parallel to the mountain belt [Faure, 1995]. Before the widespread post-orogenic Permian extension, the Stephanian period represents a transition with numerous strike-slip faults and associated intracontinental coal-bearing basins. In the French Massif central, the Sillon Houiller is one of these strike-slip faults (Fig. 1), coeval to the formation of numerous deep and narrow Stephanian basins first infilled with potassic volcanism [Letourneur, 1953]. Potassic volcanism is also known in other Stephanian basins, as for example along the Argentat fault in the western part of the Massif central (Fig. 1). These two faults meet together in the southwestern Massif central, where a comparison of these late orogenic potassic volcanism in coeval strike-slip and extensional basins can be done.

## **Geological setting**

The present study focuses on the volcanism located in the Decazeville basin, at the southern termination of the Sillon Houiller, and in Figeac and Lacapelle-Marival basins along the Argentat fault.

The Sillon Houiller fault is a 270 km-long, nearly N-S to NNE-SSW striking left-lateral fault, which crosses the entire Massif central (Fig. 1). The cumulated horizontal displacement has been estimated as ranging from 70 to 100 kilometres [Grolier and Letourneur, 1968]. This fault has been considered as a transfer fault accommodating differences between NW-SE and NE-SW extension directions on its western and eastern sides [Burg *et al.*, 1990]. However, this interpretation is not supported by the age of extension, which is Namurian-Westphalian for the western NW-SE trend and Stephanian-Autunian for the NE-SW extension [Faure, 1995]. Nevertheless, due to its NNE-SSW trend and connection with coeval Stephanian basins, the Sillon Houiller fault can be considered either as a transfer [Faure, 1995] or transform fault [Basile, 2006] in Stephanian times. The formation of the Decazeville basin was tectonically controlled by its horsetail splay termination in a transtensional regime [Basile, 2006]. In the lowermost part of the basin infilling, a thick volcanic complex overlies coarse conglomerates (Fig. 1). The upper part of the 1 800 m-thick sedimentary cross-section is almost devoid of volcanism, expressed only by a few thin layers of tuffs intercalated in lacustrine to deltaic continental sediments.

The Argentat fault is a 160 km-long curved fault in the western part of the Massif central, running from the Bosmoreau Carboniferous basin to the north and the Decazeville basin to the south (Fig. 1). This fault strikes N-S in its northern part and NW-SE in its southern part. The Argentat fault first acted as a right-lateral ductile normal fault, then as a brittle strike-slip fault during the Late Carboniferous [Feix *et al.*, 1987; Roig *et al.*, 1997]. Figeac, St Perdoux and

Lacapelle-Marival represent the southernmost basins associated with this fault (Fig. 1). They are not directly bound by the Argentat fault, but appear as the Argentat basin located on a basement flexure [Genna *et al.*, 1998]. Due to of Mesozoic sedimentary cover, the connection between Figeac and Lacapelle-Marival basins is presently hidden (Fig. 1), but both basins probably belong to a single one whose remnants were drilled below the Mesozoic sediments [Lefavrais-Raymond *et al.*, 1990]. As for the Decazeville basin, the 300 to 400 m-thick volcanic complex is located at the base of the basin infilling, overlain by coarse detrital sediments. Numerous NW-SE-trending feeder dikes also cross-cut the basement close to the Lacapelle-Marival basin (Fig. 1), providing evidence for a NE-SW extension during volcanism and basin subsidence.

In these basins, sediments are middle Stephanian (Stephanian B) in age, with early Stephanian ages (A) suspected in the St Perdoux basin, and the base of late Stephanian (C) observed in the Decazeville basin (Fig. 1) [Vetter, 1968]. Permian sediments unconformably overlay Stephanian beds in the eastern part of the Decazeville basin. They were also found in the Figeac basin [Lefavrais-Raymond *et al.*, 1990]. The age of volcanism is also assumed to be Stephanian, but this age is controversial in the Decazeville basin. From palynology, the older sediments were dated as the base of middle Stephanian [Vetter, 1968], whereas U-Pb dating on unlocated volcanic zircons provided an older Visean age [333 Ma, Bruguier *et al.*, 1998].

Volcanism is expressed as basalts, intermediate or felsic rocks, mainly as lava flows, but also as ejecta and numerous dikes in the crystalline basement. Morre [1966] and Morre and Quesnel [1967] emphasized the potassium-rich character of these lava, especially in intermediate rocks in which K<sub>2</sub>O ranges from 6 to 11%.

- In the northern part of Decazeville basin, samples were collected from two areas (Fig. 1), the Bouran section along the D963 road (samples CLB95-1 to -8), and the Lot river

banks on either side of Livinhac bridge (CLB95-9 to -17, and CLB96-1). The Bourran section consists of volcanoclastic rocks and volcanic flows conformably overlain by detrital sediments [Campagnac unit, Vetter, 1968]. The graded bedded volcanoclastic rocks rest unconformably on Variscan micaschists, and consist of breccias and lapilli-tuffs, interlayered with massive volcanic flows. The breccias are composed of meter-sized volcanic blocks caught in a tuffaceous matrix while the lapilli-tuffs include centimeter-sized angular volcanic fragments. The lapilli-tuffs locally grade to millimeter-sized crystal tuffs. Near Livinhac, massive basalts are exposed on both sides of the Lot river. The northern outcrop is cross-cutted by a porphyric dike (CLB95-17).

- In the Figeac area, two sections were sampled, East (Célé) and North (Planioles) of the city, respectively (Fig. 1). Along the Planioles section, hyalopilitic basalts crop out (CLB96-3). They are successively overlain by vitric tuffs, very altered rhyolites (CLB96-2) and detrital sediments. In the Célé valley, two outcrops of massive flows of intermediate composition (CLB96-5) can be found.

- Near Lacapelle-Marival, two volcanic levels are interbedded with Carboniferous shales, sandstones and conglomerates. Sampling concerns massive flows of mafic to intermediate composition in the northern part of the basin (CLB96-13 to -17) and small outcrops of intermediate to acidic lavas to the south (CLB96-6 to -12, Fig. 1).

### **Petrography and mineralogy**

The studied volcanic rocks are frequently severely weathered. Plagioclase is often replaced by albite or sericite, or even quartz and epidote. Olivine is transformed to iddingsite, serpentine or smectites. Orthopyroxene and clinopyroxene remain more commonly fresh. Biotite and hornblende are often chloritized. When altered, Ti-rich oxides are filled with

titanite. Mafic glass has crystallized into chlorite  $\pm$  smectites while vesicles are filled with calcite or smectites. The groundmass of intermediate lavas and rhyolites consists of fine grained quartz associated with alkali feldspar (albite or adularia). As no metamorphism has been observed in the sediments overlying the lavas, the occurrence of chlorite probably indicates hydrothermal alteration coeval with lava eruption.

In both Decazeville and Lacapelle-Marival basins, mafic rocks are commonly glomeroporphyritic with clots of large zoned plagioclase, orthopyroxene and clinopyroxene. These phenocrysts are associated with either olivine or amphibole and biotite  $\pm$  quartz. The groundmass includes plagioclase microliths and oxides. Quartz xenocrysts rimmed by orthopyroxene may occur. In Lacapelle-Marival area, some basalts differ by their trachytoid texture that consists of plagioclase laths (30 %) and less abundant larger phenocrysts (~ 10 %). The intermediate lavas exhibit seriate textures with millimeter-sized alkali feldspar phenocrysts. Their groundmass includes sandstone inclusions and quartz-filled flattened vesicles. The rhyolites display quartz and feldspar phenocrysts (20 %) and biotite needles (~ 10 %), that includes apatite and zircon. The Figeac rhyolites differ from those from Lacapelle-Marival by the presence of green hornblende.

When preserved (CLB95-5, CLB95-16, CLB96-1, CLB96-3), plagioclase composition of mafic volcanic rocks varies from Ca-rich cores (An<sub>98-60</sub>) to Na-rich rims (An<sub>50-40</sub>). Preserved alkali feldspar (CLB96-11) gives sanidine compositions (Ab<sub>57-66</sub>). Clinopyroxene phenocrysts (CLB96-1, CLB95-16, CLB95-5) show homogeneous augite compositions [Wo<sub>39-44</sub> En<sub>44-52</sub> Fs<sub>12-21</sub>; Morimoto *et al.* 1988], while preserved orthopyroxene (CLB95-5, CLB95-16) has bronzite compositions (Wo<sub>2-2.5</sub> En<sub>77-78</sub> Fs<sub>19-21</sub>).



## Geochemistry

### Analytical techniques

Thirty one samples have been analysed for whole rock major- and trace-element concentrations. Among them, twenty one have been analysed for Nd, and nine for Sr isotopic chemistry (tabl. I). Major and compatible trace-elements have been measured by ICP-AES at the Centre de Recherches Pétrographiques et Géochimiques de Nancy (samples CLB95) or by X-ray fluorescence (XRF) at the Earth Sciences Department of the Australian National University at Canberra (Australia) (samples CLB96). The trace-element concentrations have been determined by inductively coupled plasma-mass spectrometry (ICP-MS, VG-PQ<sup>2+</sup>) at the Laboratoire de Géodynamique des Chaînes Alpines de l'Université Joseph Fourier (Grenoble), after acid dissolution, using the procedures described by Barrat *et al.* [1996]. Standards used for the analyses were JB2, WSE, BIR-1, JR1, and BHVO.

Sr and Nd isotopic ratios were measured at the Laboratoire de Géochemie isotopique de l'Université Paul Sabatier (Toulouse), on a Finnigan MAT261 multicollector mass spectrometer using the analytical procedures of Lapierre *et al.* [1997]. Results on La Jolla Nd standards yielded  $^{143}\text{Nd}/^{144}\text{Nd} = 0.511850 \pm 8$  (mean on 39 runs). Results on NBS 987 Sr standard yielded  $^{87}\text{Sr}/^{86}\text{Sr} = 0.710250 \pm 5$  (mean on 200 runs). The isotopic data have been corrected for “in situ decay” for an age of 300 Ma.

### Alteration and elemental mobility

Most of the analysed rocks have a loss on ignition lower than 6% (LOI, tabl. I), due to the presence of calcite- or smectites-filled vesicles. Moreover, the altered nature of the studied Carboniferous volcanic rocks means that before any petrological inferences can be drawn from their chemistry, possible effects of subsolidus transformations must be considered.

As Zirconium (Zr) is considered to be immobile during low-grade alteration of igneous rocks of mafic to felsic composition by hydrothermal fluids [Gibson *et al.*, 1982], it has been used to test the mobility of minor ( $\text{TiO}_2$ ), and incompatible trace elements (Rb, Sr, Pb, La, Nb, Y, U and Th) in figure 2. Rb, Sr, Pb and La do not display any correlation with Zr. This implies that Large Ion Lithophile Elements (LILE) have been extensively mobilised. For example, some samples are enriched in Pb (others in Ba), probably in relation with hydrothermal circulations [Marignac and Cuney, 1999]. Thus, variations in LILE are not expected to reflect the composition of the sources (Sr isotopic ratios) or the magmatic processes. These variations will not be discussed further on. Similarly, the behaviour of K and Ca also exclude their use for sample classification.

In contrast, other elements such as U, Th, Nb or  $\text{TiO}_2$  correlate rather well with Zr, suggesting relative immobility. However, some samples (CLB95-17, 95-15, 95-1, 96-5) differ from the correlation trend on the Nb-Zr plot. The same samples, together with CLB95-6, 95-12, 96-2, 96-8 show a negative correlation on the  $\text{TiO}_2$ -Zr diagram, while all other samples exhibit a positive correlation. CLB95-17 is the most mafic sample ( $\text{SiO}_2$  51.9%, MgO 7.7%, tabl. I); CLB95-1, 95-6, 95-12, 95-15, 96-2, 96-5, 96-8 represent the most  $\text{SiO}_2$ -rich and MgO-poor samples ( $\text{SiO}_2 > 65\%$  and  $\text{MgO} < 1.3\%$ , tabl. I).

In the Y-Zr diagram, each sampling site corresponds to distinct Y-Zr correlation trends. This suggests that the Carboniferous volcanic rocks are not related by simple fractionation processes, but rather reflect variations of partial melting ratios, and/or complex fractionation processes dominated by clinopyroxene or amphibole accumulation and/or removal.

### Volcanic rocks terminology

Because of the alteration experienced by the Carboniferous volcanic rocks, it was difficult to name the different facies on the basis of mineralogy only, especially for the mafic rocks.

The various types of volcanic rocks have been discriminated using Ti and immobile trace elements Zr, Nb and Y (figure 3). The most mafic rock (CLB95-17) plots in the andesite/basalt field. The samples that exhibit a positive correlation in the  $\text{TiO}_2$ -Zr diagram (figure 2) have a constant Zr/Ti ratio and plot near the fence separating the field of rhyodacites from that of andesites, with the exception of sample 95-13 which plots in the alkali basalt field. Among the most  $\text{SiO}_2$ -rich samples that show a negative correlation in the  $\text{TiO}_2$ -Zr diagram (figure 2), CLB95-1 plots in the trachy-andesite field, CLB95-6, 95-12, 96-2 and 96-8 in the rhyodacite field, CLB96-5 in the rhyolite field and CLB95-15 at the fence between trachyte and comendite.

Because K is highly mobile during alteration and low grade metamorphism processes, we cannot rely on the K contents of the Carboniferous volcanic rocks (half of the samples have K contents from 6 to 12%) to infer calc-alkaline or shoshonitic affinities. Müller *et al.* [1992] proposed a Th/Yb versus Ta/Yb correlation diagram to discriminate arc-tholeiites, calc-alkaline and shoshonitic suites. As shown in figure 4, most of the Carboniferous volcanic rocks fall in the shoshonitic field, with the exception of 95-17 andesite/basalt which fall in the calc-alkaline field. In this plot the most  $\text{SiO}_2$ -rich samples exhibit a more shoshonitic affinity.

#### Trace-element geochemistry

Figure 5 illustrates the chondrite-normalized [Sun and McDonough, 1989] Rare Earth Elements (REE) patterns of the Carboniferous volcanic rocks. All the rocks are enriched in Light (L) REE relative to the heavy (H) REE [ $2.8 \leq (\text{La/Yb})_n \leq 31.9$ ]. However, HREE patterns are either flat or gently sloping, suggesting that LREE enrichment reflects a metasomatised source [Keppler, 1996; Tatsumi and Kogiso, 1997; Beccaluva *et al.*, 2004]. All samples also show negative Eu anomalies ( $0.38 < \text{Eu}/\text{Eu}^* < 0.84$ ). These anomalies suggest an early fractionation of plagioclase during magma differentiation. The most  $\text{SiO}_2$ -rich rocks

(e.g. samples CLB95-1 and -6 in Bourran) are commonly less enriched than mafic rocks (e.g. samples CLB95-4 and -8 in Bourran), especially for HREE (Figure 5). This suggests again that these lavas cannot result from fractional crystallisation of basalts.

There are no significative differences between the sampling sites, with the exception of negative Ce anomalies ( $0.67 \leq \text{Ce}/\text{Ce}^* \leq 0.38$ ) for four rhyodacites from the southern part of Lacapelle-Marival basin (CLB96-6, CLB96-10 to -12). Two rhyodacites from Decazeville (CLB95-8, CLB95-9) differ from the other rocks by flatter patterns [ $2.8 < (\text{La}/\text{Yb})_n < 4.9$ ]. The trachyte from the Livinhac site (CLB95-15) differs from the other felsic rocks by a very steep REE pattern due to an important HREE depletion [ $(\text{La}/\text{Yb})_n = 31.9$ ].

Figure 6 shows the primitive mantle-normalized spiderdiagrams [Sun and McDonough, 1989]. Whatever their location or mafic character, Carboniferous volcanic rocks exhibit similar trace element patterns, characterized by negative Nb, Ta, P, Sm, Ti and positive Th, U and Pb anomalies, respectively. As shown previously (Fig. 2), Sr vary from 10 to 330 ppm. The most mafic rocks exhibit a homogeneous composition for all trace elements, with the exception of CLB95-17 that differs from the other mafic rocks by a Zr and Hf depletion. CLB95-17 as other samples (CLB95-1, 95-13, 95-15, 96-5) that define a negative trend in  $\text{TiO}_2/\text{Zr}$  correlation (Fig. 2) are all characterized by a negative Y anomaly.

#### Nd and Sr isotopic geochemistry

The Carboniferous volcanic rocks span a rather small range of  $\epsilon_{\text{Nd}}$  values, from -5.8 to -3.9, with the relative exception of CLB96-5 rhyolite ( $\epsilon_{\text{Nd}} = -2.2$ ). All  $\epsilon_{\text{Nd}}$  are negative, and do not present significative variations with rock type or rock chemistry (tabl. I). These very homogeneous values suggest that these lavas are cogenetic, and indicate a crustal signature in the melting source. The  $^{87}\text{Sr}/^{86}\text{Sr}$  ratio is less homogeneous and ranges between 0.71128 and 0.70415;  $\epsilon_{\text{Sr}}$  ranges from 0 to 325 (tabl. I). As exposed above, Rb and Sr mobility during

hydrothermalism and alteration precludes any use of these elements as markers of the magmatic sources.

## **Discussion**

### Geochemistry

Whatever their location or petrology, all the analysed samples display high  $(La/Yb)_n$  ratios but almost flat HREE patterns, marked negative Eu anomalies, homogeneous negative  $\epsilon_{Nd}$  values (between  $-5.8$  and  $-3.9$ ) and in primitive mantle-normalized plots, negative Nb, Ta, P, Sm, Ti anomalies and positive Th, U and Pb anomalies. The most  $SiO_2$ -rich and MgO-poor rocks and one basalt (CLB95-17) can be distinguished from other samples by a negative correlation in Ti/Zr diagram (figs. 2 & 3). These samples present additional geochemical characteristics, such as Y depletion and enhanced negative Ti anomalies, and for some samples negative Ce anomalies.

The occurrence of mafic rocks implies partial melting of the mantle. As the most  $SiO_2$ -rich rocks display REE spectrums that are less enriched in LREE than mafic rocks, they can not be linked together only by crystal fractionation processes. However, as they present similar characteristics, all these lavas may have been formed from similar sources and similar mechanisms. All these rocks belong to a shoshonitic suite, characterized by the absence of Ti and Fe enrichment during crystal fractionation. This is a classical feature of hydrous melts [Gill, 1981]. Similarly, as Nb and Ta are insoluble in water, their negative anomalies in primitive mantle-normalized plots can also result from a mantle source metasomatised by water.

As shown by the occurrence of sedimentary xenoliths in these lavas, the positive U, Th and Pb anomalies may be explained by contamination by continental crust, while the negative Ce

anomaly may indicate contamination by oceanic rocks (sediments or altered oceanic crust) [Bohrson and Reid, 1997]. These contaminations may occur during magma ascent, but more probably reflect the direct contribution of hydrous crustal materials to mantle melting, as suggested by the negative  $\epsilon_{\text{Nd}}$  values, the homogeneous shape of REE patterns, and the Ti/Y versus Zr/Y and Nb/La versus Th/La diagrams (Fig. 7). All samples display a positive linear trend, except one basalt (CLB95-17) in both diagrams and the most SiO<sub>2</sub>-rich rocks in Ti/Y versus Zr/Y diagram. This suggests that the contamination by hydrous crustal material is strictly linked to melting of the source, i.e. that the mantle protolith was metasomatised by hydrous crustal materials, that participated in its melting. Among the analysed rocks, the Decazeville acidic lavas have the highest Ti/Y, Zr/Y, Nb/La and Th/La ratios, suggesting that the proportion of crustal material was higher in the source of these volcanic rocks than for Figeac and Lacapelle-Marival magmas.

Initial fractionation of apatite from these melts can explain the P negative anomalies. Similarly, the negative anomalies in Ti, Eu and Y can be explained by early fractionation of Fe-Ti oxides (sometimes found as inclusions in the phenocrysts), plagioclase and clinopyroxenes, respectively. The peculiar Y depletion in the SiO<sub>2</sub>-rich lavas and CLB95-17 basalt can be explained by clinopyroxene or amphibole fractionation, that may have not occurred for the other lavas. Similarly, their negative Ti/Zr correlation can indicate Fe-Ti oxide fractionation.

#### Structural control on magmatism

Despite differences in structural setting, there are no marked differences in the petrology or geochemistry of rocks from Decazeville, Figeac and Lacapelle-Marival basins. Thus, the faults and the tectonic regime control the formation of these Stephanian basins, where the

volcanism took place, but they do not influence its petrology or geochemistry. The NW-SE striking dikes found in the basement of Lacapelle-Marival basin, and the occurrence of volcanism as the first infilling of all the basins along the Sillon Houiller fault indicate a strike-slip tectonic regime at the French Massif central scale, with a NE-SW stretching direction and a NW-SE shortening direction. This regime is associated to the left-lateral slip on the Sillon Houiller fault, and to extension or right-lateral transtension along the Argentat fault. This seems to be the main tectonic regime at the time of the infilling of the Stephanian basins along these faults [Basile, 2006]. As the same volcanism systematically occurred in all the basins along the Sillon Houiller fault [Letourneur, 1953], it can be proposed that the Lacapelle-Marival volcanism, located only 25 km West of the Sillon Houiller fault, came from the same source as the Decazeville volcanism. The local variations in the tectonic regime only controlled the volume and surface covered by volcanic lava flows.

#### Late-orogenic Variscan magmatism

Stephanian volcanism from the French Massif central belongs to the extended K-rich igneous province emplaced during Stephanian to Early Permian times in the southern part of the European Variscan belt [Pyrénées: Innocent *et al.*, 1994; Western Alps: Cannic *et al.*, 2002; Central Alps: Schaltegger *et al.*, 1991; Southern Alps: Rottura *et al.*, 1998, Cortesogno *et al.*, 1998; Corsica and Sardinia: Tommasini *et al.*, 1995]. In the Pyrénées [Bixel and Lucas, 1983] and Western Alps [Capuzzo and Bussy, 2000], this volcanism is located in basins clearly controlled by strike-slip faults. Together with the K-rich character, these igneous provinces share similar geochemical patterns with the Stephanian volcanism for the SW Massif central: in chondrite-normalized diagrams, LREE-enriched patterns, flat HREE spectra and negative Eu anomalies; in primitive mantle normalized diagrams, negative Nb, Ta and Ti anomalies, and positive Th and U anomalies; and negative  $\epsilon_{Nd}$  values (-3 to -7), with the

exception of the most recent (Permian) lava that have slightly positive  $\epsilon_{\text{Nd}}$  values [Innocent *et al.*, 1994; Rottura *et al.*, 1998; Cannic *et al.*, 2002;]. It is noteworthy that these geochemical characteristics were homogeneous during a large time span and in various parts of the Variscan belt.

The origin of potassic late-orogenic magmatism is still a matter of debate [Gibson *et al.*, 1993; Liégeois *et al.*, 1998; Miller *et al.*, 1999; Pecerrillo and Martinotti, 2006].

- As this magmatism indicates a hydrous melt, some authors [e.g. Stampfli, 1996] proposed that it was generated in an active subduction or back-arc setting. However no subduction was active in this part of the Variscan belt at that time, the studied Stephanian volcanism can therefore not be directly linked to subduction processes.

- More frequently, the geochemical patterns of potassic late-orogenic magmatism have been explained by contamination, either by crustal contamination during magma ascent, or more probably by mantle metasomatism before melting, as indicated by the systematic and homogeneous character of contamination. Up to now, this metasomatism has been considered to be inherited from a suprasubduction position, either in the same orogenic cycle, or during previous subductions [e.g. Miller *et al.*, 1999].

The southern part of the Variscan orogeny results from continental collision following a northward subduction, whose remnants are found in the French Massif central [Matte, 1986, 2001]. However, the southernmost part of the French Massif central as well as the Pyrénées belong to the subducting plate, and its underlying mantle was not in a suprasubduction position, and could not be metasomatised by a Variscan subduction. However, this does not exclude older (Precambrian) metasomatism processes.

If the mantle was previously metasomatised, the main mechanism proposed to explain its melting in a post-orogenic setting is lithospheric thinning, inducing mantle decompression [e.g. Gibson *et al.*, 1993; Innocent *et al.*, 1994; Rottura *et al.*, 1998]. However, the Stephanian



volcanism occurred in a strike-slip tectonic regime in the French Massif central, and before the main Permian extension that ended the variscan orogenic cycle [Ménard and Molnar, 1988].

- We propose an alternative mechanism, controlled by lithospheric-scale strike-slip faulting, to explain both mantle metasomatism and mantle melting generating the potassic late-orogenic volcanism. At the end of the orogeny, the various parts of the mountain belt differ by their crustal and lithospheric thicknesses, thermal gradients, rock nature and geological history. As observed in the Ivrea body, hydrous metasediments can occur in the lower crust at the end of the Variscan orogeny [Harlov and Forster, 2002].

As the late-orogenic strike-slip faults cross the orogenic zonation with some tens or hundreds kilometres of horizontal slipping, these bring into contact different parts of the mountain belt. For example, this horizontal displacement can bring into contact, and at the same depth, mantle materials on one side and lower crust, including still hydrated metasediments, on the other. The crustal fluids could then locally hydrate the adjacent mantle, and this metasomatism may induce mantle melting. The resulting magmas have geochemical patterns that indicate contamination by continental crust.

In this hypothetical mechanism, mantle metasomatism is not inherited from a previous suprasubduction zone, but is acquired by horizontal fluid transfer in or across lithospheric-scale strike-slip faults. Consequently, mantle melting can occur without vertical displacements and adiabatic decompression, since metasomatism controlled by strike-slip faults drastically decreases its solidus temperature. Because fluid transfers across strike-slip faults and induced mantle metasomatism are only local and can occur only once, this mechanism can explain why this late-orogenic potassic magmatism represents small volumes of lava, and why it happens only at the beginning of late-orogenic strike-slip faulting.

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## Figure Captions

Figure 1: Schematic geologic map of the area between Lacapelle-Marival and Decazeville basins. Analysed samples are located. A: Schematic geologic map of the French Massif central [from Basile, 2006]. B: Synthetic lithostratigraphic column of the Decazeville basin [from Basile, 2006].

*Figure 1: Carte géologique schématique de la région comprise entre les bassins de Lacapelle-Marival et Decazeville, avec la localisation des échantillons analysés. A: Carte géologique schématique du Massif central français [d'après Basile, 2006]. B: Colonne lithostratigraphique synthétique du bassin de Decazeville [d'après Basile, 2006].*

Figure 2: Variations of minor- ( $\text{TiO}_2$ ) and trace-elements (Th, U, Nb, La, Pb, Rb, Sr, Y) versus Zr. The symbols indicate the sample location (cf. Fig. 1).

*Figure 2: Variations des teneurs en éléments mineur ( $\text{TiO}_2$ ) et traces (Th, U, Nb, La, Pb, Rb, Sr, Y) par rapport au Zr. Les symboles correspondent à la localité d'échantillonnage (cf. Fig. 1).*

Figure 3: Zr/Ti versus Nb/Y discrimination diagram [Winchester and Floyd, 1977].

*Figure 3: Diagramme discriminant Zr/Ti par rapport à Nb/Y [Winchester and Floyd, 1977].*

Figure 4: Th/Yb versus Ta/Yb discrimination diagram [Müller *et al.*, 1992].

*Figure 4: Diagramme discriminant Th/Yb par rapport à Ta/Yb [Müller *et al.*, 1992].*

Figure 5: Chondrite-normalized rare earth elements patterns [Sun and McDonough, 1989]. Sample sets as a function of location (cf. Fig. 1).

*Figure 5: Spectres de terres rares normalisés aux chondrites [Sun and McDonough, 1989]. Les échantillons sont regroupés en fonction de leur localisation (cf. Fig. 1).*



Figure 6: Primitive mantle-normalized spidergrams [Sun and McDonough, 1989]. Sample sets as a function of location (cf. Fig. 1).

*Figure 6: Spidergrams normalisés au manteau primitif [Sun and McDonough, 1989]. Les échantillons sont regroupés en fonction de leur localisation (cf. Fig. 1).*

Figure 7: Variations of Ti/Y versus Zr/Y (left) and Nb/La versus Th/La (right). The symbols indicate the sample location (cf. Fig. 1).

*Figure 7: Variations des rapports Ti/Y en fonction de Zr/Y (à gauche) et Nb/La en fonction de Th/La (à droite). Les symboles indiquent la localisation des échantillons (cf. Fig. 1).*

## Table Captions

Table I: Major- and trace-element analyses,  $^{87}\text{Sr}/^{86}\text{Sr}$  and  $^{143}\text{Nd}/^{144}\text{Nd}$  isotope ratios for studied samples. Major oxides are in wt%, Trace-elements are in ppm.  $\text{Fe}_2\text{O}_3$ : total iron as ferric oxide; LOI: loss on ignition; bdl: below detection limit; nd: not determined. Rock types are given from figure 3. Location on figure 1 (LM for Lacapelle-Marival).

*Table I: Analyses chimiques en éléments majeurs et traces, rapports isotopiques en  $^{87}\text{Sr}/^{86}\text{Sr}$  et  $^{143}\text{Nd}/^{144}\text{Nd}$  pour les échantillons étudiés. Les éléments majeurs sont donnés en %, les éléments traces en ppm.  $\text{Fe}_2\text{O}_3$ : Fer total; LOI: perte au feu; bdl: sous le seuil de detection; nd: non déterminé. La terminologie des laves est déduite de la figure 3. Localisation sur la figure 1 (LM: Lacapelle-Marival).*

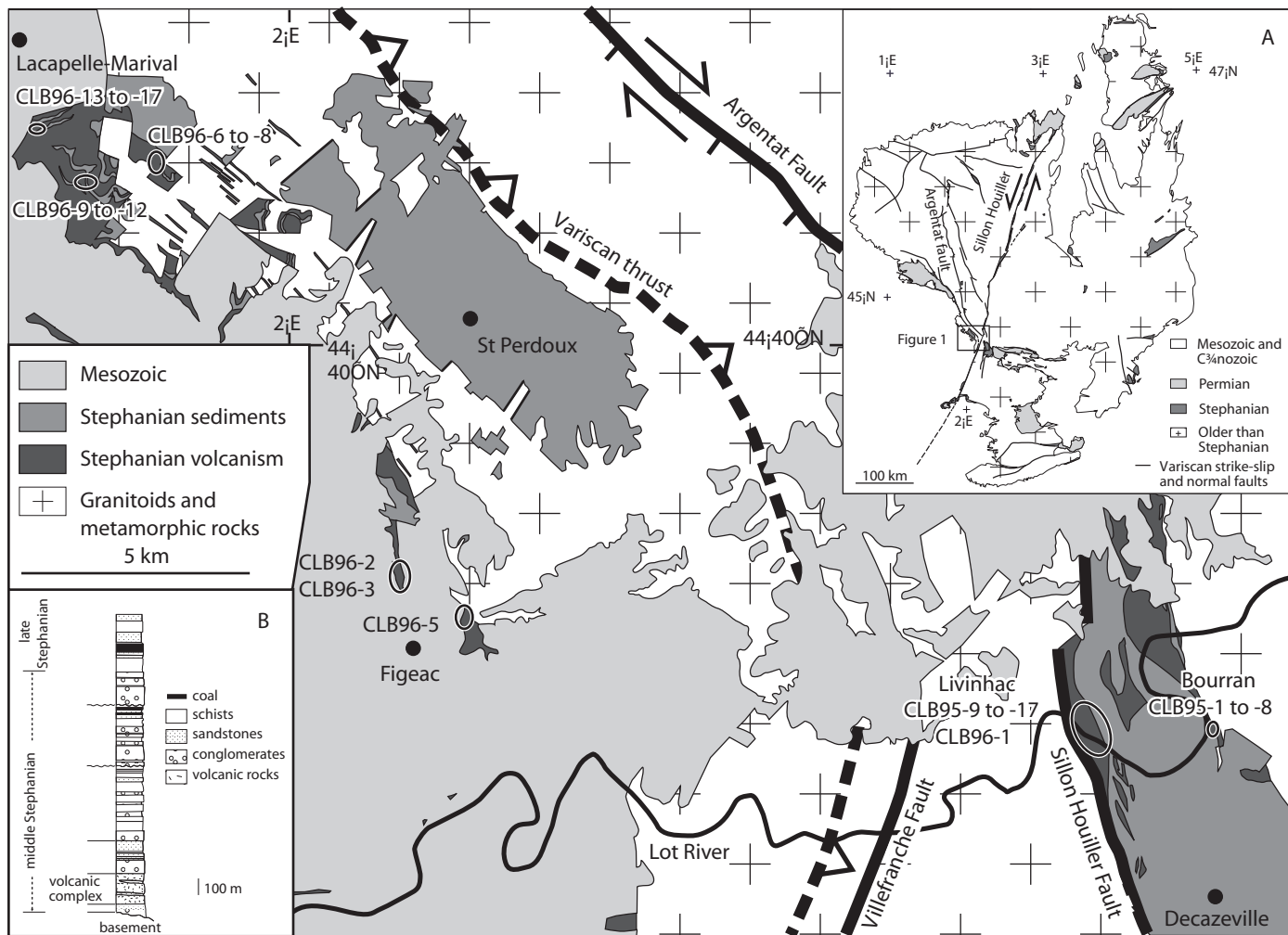


Figure 2

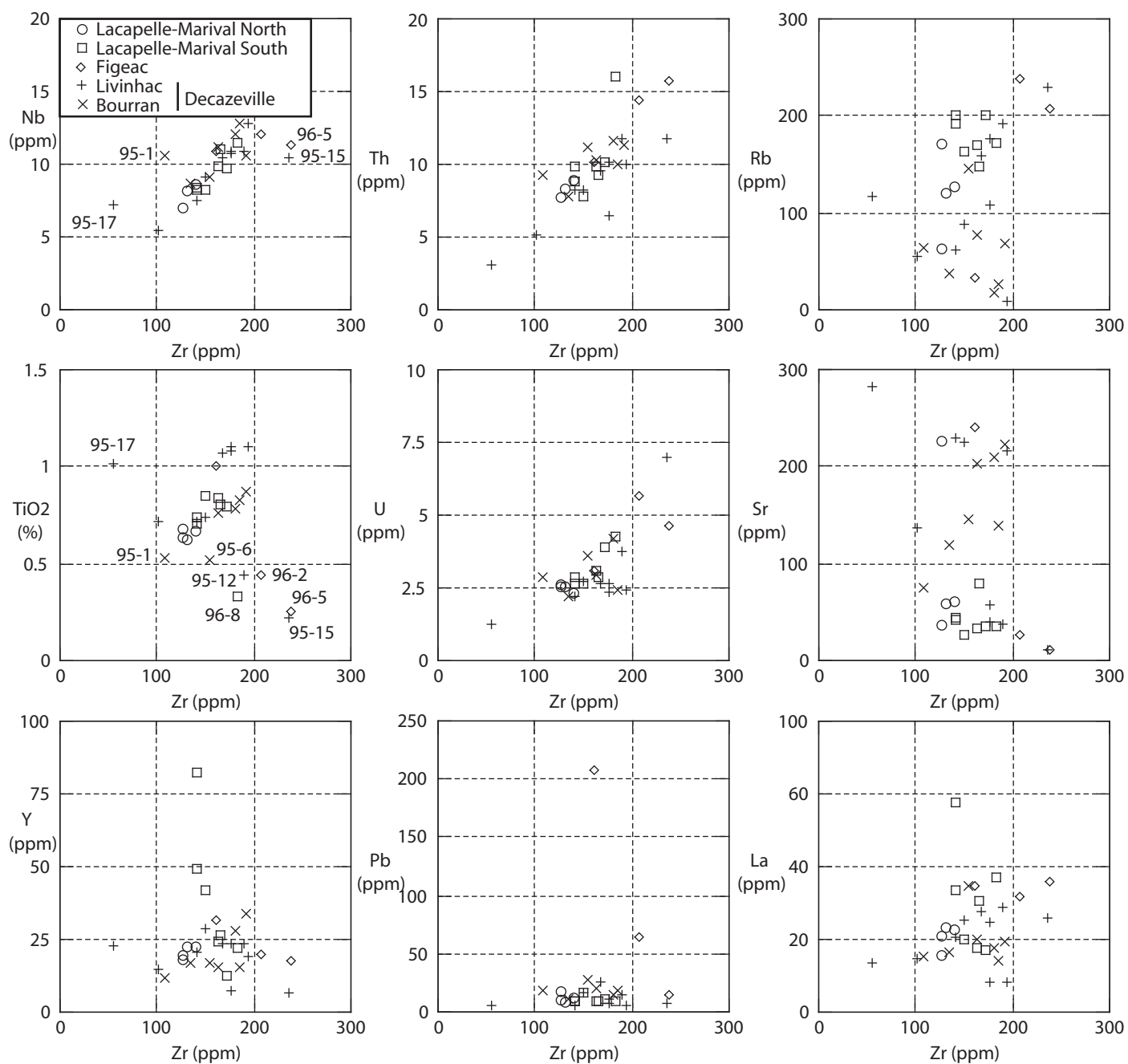


Figure 3

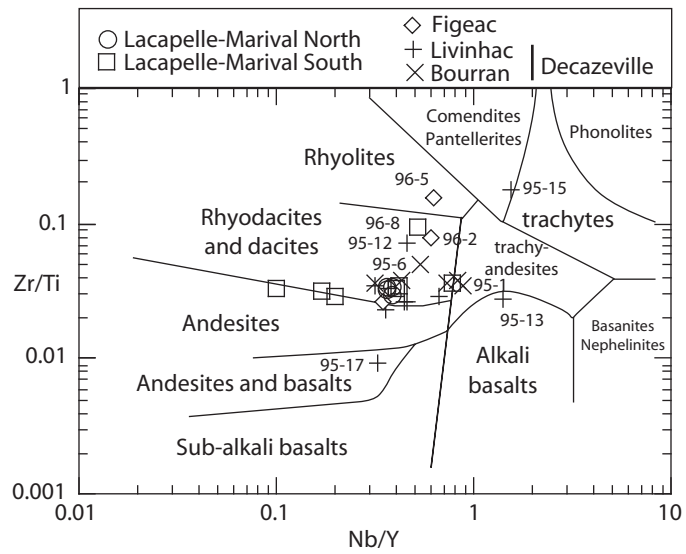


Figure 4

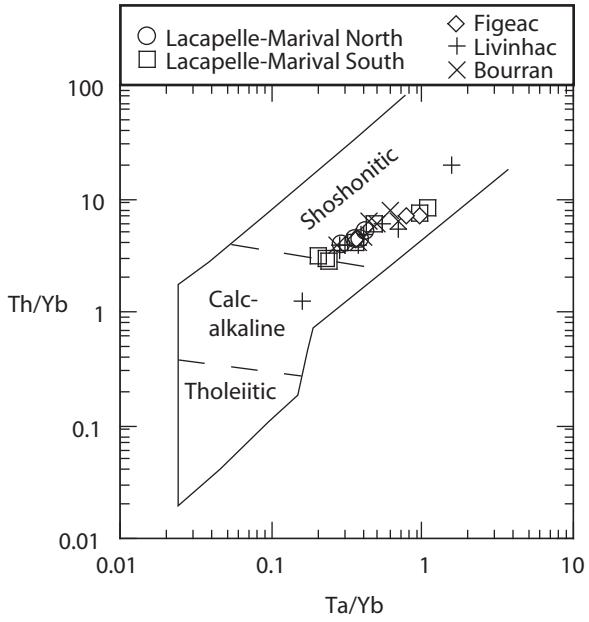


Figure 5

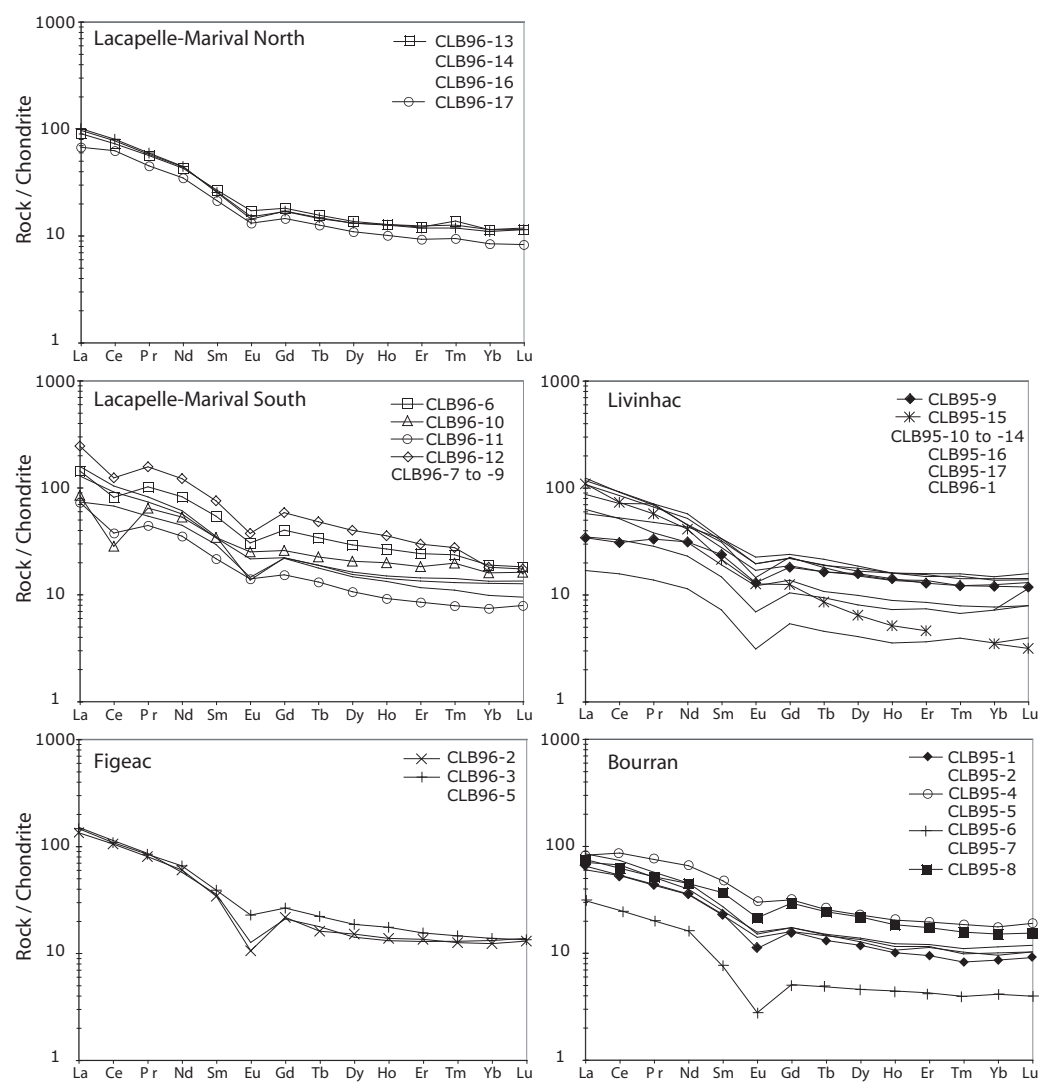


Figure 6

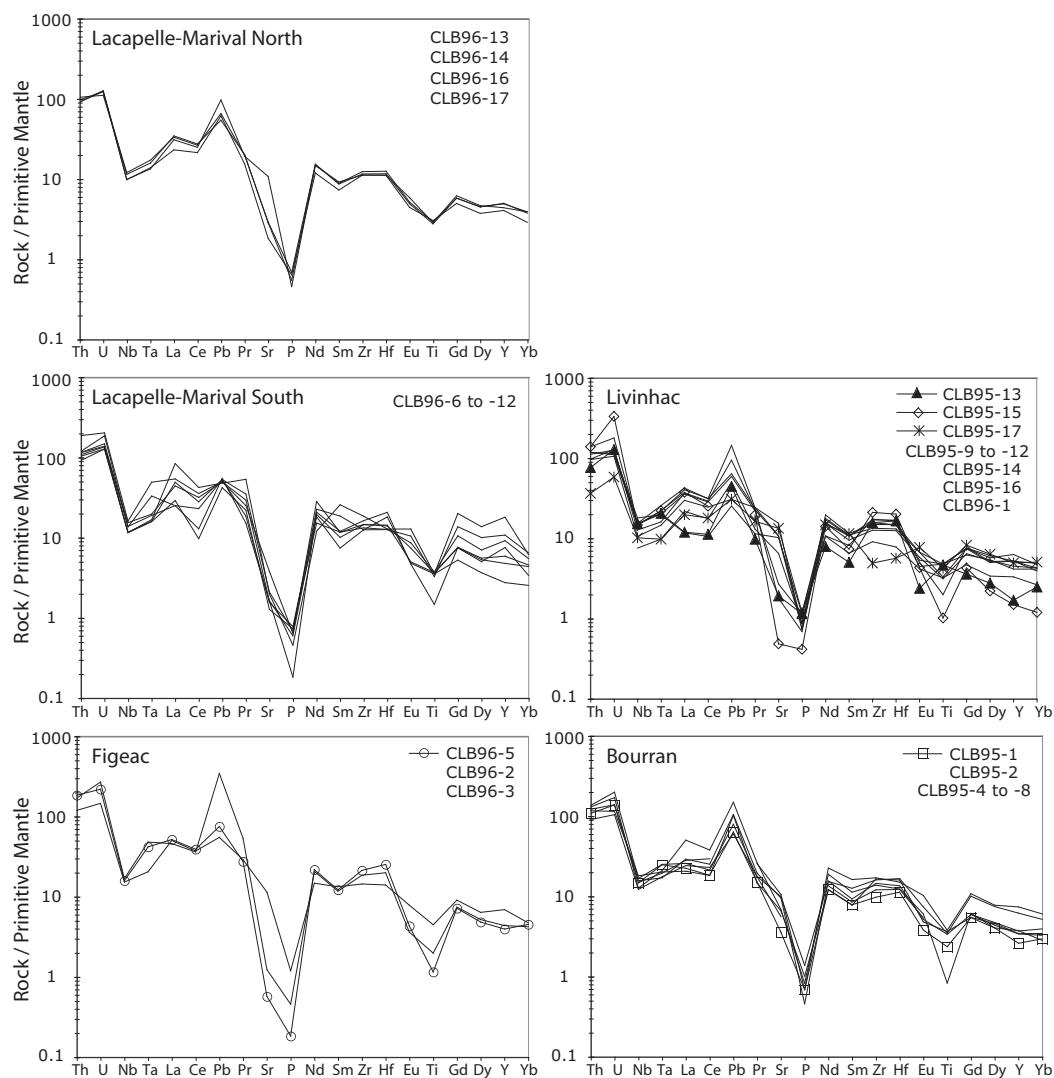
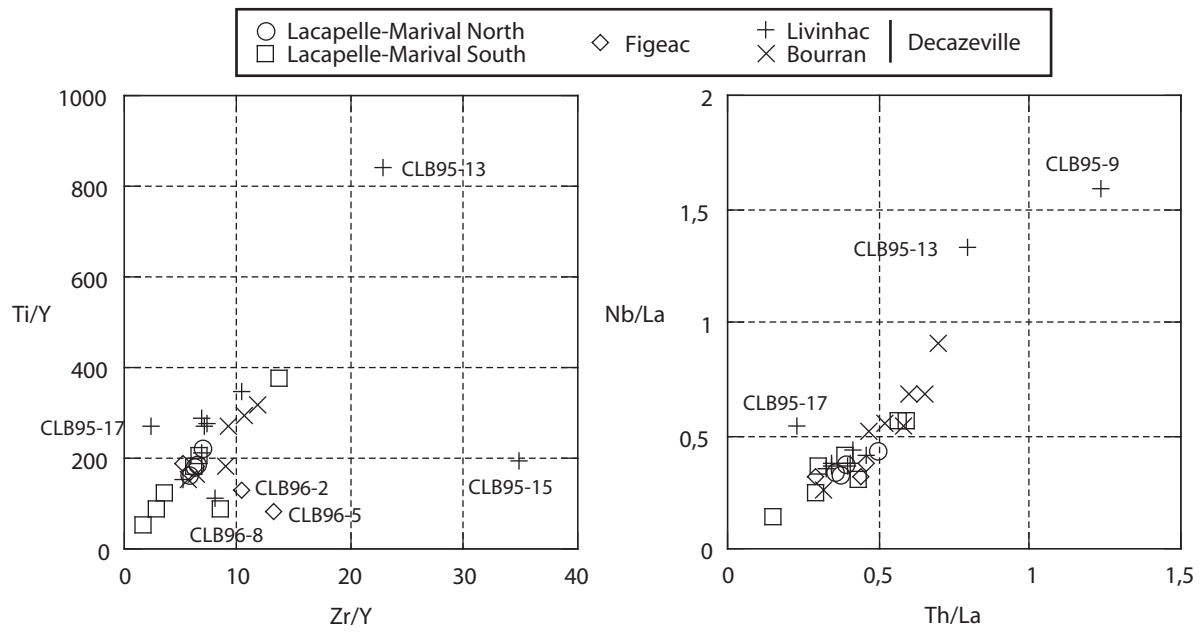


Figure 7





Location Sample Number Rock Type	Bourran CLB95-1 trachy-andesite	Bourran CLB95-2 rhyodacite	Bourran CLB95-4 rhyodacite	Bourran CLB95-5 rhyodacite	Bourran CLB95-6 rhyodacite	Bourran CLB95-7 rhyodacite	Bourran CLB95-8 rhyodacite	Livinghac CLB95-9 rhyodacite	Livinghac CLB95-10 rhyodacite	Livinghac CLB95-11 rhyodacite	Livinghac CLB95-12 rhyodacite	Livinghac CLB95-13 alkaline basalt	Livinghac CLB95-14 rhyodacite	Livinghac CLB95-15 trachyte	Livinghac CLB95-16 andesite/basalt	Livinghac CLB95-17 rhyodacite	Livinghac CLB96-1 rhyodacite
SiO <sub>2</sub>	65.65		55.12	61.9	68.02	58.38	59.62	59.04	56.97	59.44	67.98	59.99	58.18	73.16	58.5	51.9	59.36
Al <sub>2</sub> O <sub>3</sub>	16.4		20.8	17.49	15	21.33	20.7	19.85	17.4	18.25	16.34	17.94	17.18	14.12	17.1	18.35	16.53
Fe <sub>2</sub> O <sub>3</sub>	4.48		7.05	5.29	5.65	3.96	3.4	5.87	7.08	6.5	4.32	6.4	6.78	0.9	6.89	9.16	6.82
MnO	0.06		0.15	0.13	bdl	0.07	0.07	0.07	0.08	0.07	bdl	0.09	0.09	bdl	0.07	0.13	0.08
MgO	0.77		5.69	3.76	1.06	4.53	4.21	4.35	6.64	2.92	1.29	3.47	4.66	bdl	4.83	7.69	5.12
CaO	3.98		3.37	5.49	2.14	2.46	2.68	0.98	6	1.56	0.45	2.02	7.1	bdl	6.29	5.83	6
Na <sub>2</sub> O	3.46		5.64	2.55	2.98	5.92	7.23	8.11	3.2	0.7	2.08	0.7	2.97	0.49	3	2.02	2.75
K <sub>2</sub> O	4.49		1.15	2.41	4.42	2.3	1	0.39	1.75	8.14	7.1	8.1	1.81	11.61	2.48	3.84	2.38
TiO <sub>2</sub>	0.51		0.82	0.73	0.52	0.79	0.75	1.04	0.68	1.03	0.43	1.01	1.04	0.22	0.7	0.92	0.72
P <sub>2</sub> O <sub>5</sub>	0.15		0.15	0.1	0.18	0.22	0.3	0.25	0.16	0.22	0.15	0.22	0.16	0.09	0.19	0.26	0.19
LOI	4.5		6.14	3.57	0.18	4.85	3.81	5.66	5.18	6.32	3.12	6.65	3.02	0.83	2.98	8.99	2.47
Cr	26.12	371.1		366.3	47.24	295.53	283.9	75.77	352	250.2	23.5	270.6	251.5			161.1	301
V	33.3	133.02		133.5	49.23	67.05	74.44	82.9	146.7	116.3	364.5	123	142.5			241.7	153
Sc	13.46	24.46		23.84	13.34	22.56	20.7	24.54	35.76	21.63	8.32	18.56	26.9			35.9	20
Ni	8.5	83.9		29.13	23.79	58.56	45.2	12.1	115.6	8.2	102	105.1				24.9	40
Co	3.83	22.83		5.15	20.8		19.5	18.1	29.04	22.34	5.56	22.9	24.78	1.49		22.9	22.99
Cu	7.9	25.41		9.21	11.05	63.2	9.34	21.96	40.84	24.6	4.3	27.9	32.55			23.65	18
Rb	64.43	36.7	69	77.8	146.25	26.22	17.53	8.75	61.73	175.91	192.47	108.5	158	229.07	55.7	117.9	88.47
Sr	75.11	118.43	223	202.34	144.71	139.42	210	216.3	229.7	57.31	37.82	40.6	326.6	10.2	137	282.3	226
Y	11.86	16.87	33.6	15.5	17.1	15.65	28.1	18.9	20.55	23.73	23.87	7.71	23.7	6.8	15.1	22.5	28.63
Zr	109.28	135.35	191	163.04	154.63	185.34	180.4	194.2	141.5	176.1	189.74	176.4	167	237.03	101.7	55.14	150
Nb	10.53	8.71	10.63	11.2	9.08	12.85	12	12.8	7.52	10.79	10.93	10.9	10.4	10.47	5.4	7.27	9.06
Cs	7.84	4.24	nd	2.96	6.99	5.5	4.6	4.95	2.03	22.17	19.07	9.2	53.63	3.04		21	3.4
Ba	1412	323	359	555.9	1046.7	653.7	619.6	76.05	387.3	592.73	709.07	349.9	493.7	737.82	307	503.7	508
La	15.38	16.64	19.42	20.05	34.69	14.22	17.7	8.07	20.5	24.55	29.12	8.2	27.65	25.76	14.9	13.55	25.364
Ce	32.61	40.7	52.47	44.85	67.24	32.37	37.9	18.82	43.52	50.67	55.44	19.94	55.9	44.73	31.4	31.9	51.716
Pr	4.17	4.84	7.1	5.4	7.25	4.07	4.9	3.15	6.71	6.08	6.53	2.7	6.71	5.42	3.6	4.5	6.287
Nd	16.58	18.44	30.58	21.02	25.58	16.4	20.8	14.55	20.93	23.16	21.06	10.7	26.65	19.15	14.4	20	24.095
Sm	3.5	3.78	7.22	4.14	5.02	3.45	5.6	3.63	4.22	4.77	4.74	2.24	5.11	3.27	2.9	5.11	4.873
Eu	0.64	0.91	1.73	0.87	1.03	0.81	1.24	0.75	0.98	0.9	0.84	0.4	1.13	0.73	0.7	1.3	1.134
Gd	3.25	3.54	6.45	3.55	3.67	3.26	6.01	3.71	3.84	4.43	4.59	2.14	4.51	2.56	2.8	4.9	4.48
Tb	0.49	0.56	0.95	0.55	0.55	0.54	0.91	0.61	0.61	0.68	0.67	0.35	0.7	0.32	0.4	0.8	0.697
Dy	2.99	3.49	5.75	3.23	2.99	3.4	5.52	3.99	3.9	3.7	3.8	2.04	4.5	1.63	2.5	4.7	4.211
Ho	0.57	0.69	1.15	0.6	0.58	0.65	1.04	0.8	0.78	0.8	0.77	0.41	0.9	0.29	0.5	0.9	0.889
Er	1.56	1.98	3.22	1.86	1.5	1.9	2.86	2.12	2.25	2.24	2.16	1.22	2.55	0.76	1.4	2.6	2.456
Tm	0.21	0.28	0.47	0.26	0.55	0.25	0.4	0.31	0.31	nd	nd	0.17	0.36	nd	0.2	0.4	0.38
Yb	1.46	1.94	2.98	1.62	1.4	1.7	2.55	2.04	2.11	2.1	1.93	1.22	2.4	0.59	1.3	2.5	2.321
Lu	0.23	0.3	0.48	0.26	0.24	0.26	0.39	0.3	0.33	0.31	0.29	0.2	0.36	0.08	0.2	0.4	0.356
Hf	3.5	3.77	4.67	4.2	3.81	4.97	5.15	5.11	3.86	4.56	5.22	5.09	4.6	6.21	2.42	1.75	4.06
Ta	1.02	0.72	0.8	0.7	0.86	0.83	1.03	0.81	0.6	0.83	1.05	0.84	0.9	0.91	0.4	0.4	0.66
Pb	17.6	11.12	nd	19.4	27.6	18.9	14.7	4.7	5.66	10.96	14.72	8.26	26.7	7.21	5.7	17.48	
Th	9.22	7.74	11.27	10.37	11.13	9.95	11.6	9.95	8.19	10.12	11.76	6.52	9.5	11.84	5.15	3.1	8.26
U	2.9	2.2	nd	2.91	3.58	2.42	4.2	2.42	2.22	2.32	3.75	2.68	2.65	6.97		1.22	2.7
Age: 300 Ma																	
<sup>87</sup> Sr/ <sup>86</sup> Sr			0.715103 ± 1								0.712987 ± 11		0.712618 ± 15		0.714451 ± 2		0.714744 ± 20
<sup>87</sup> Rb/ <sup>86</sup> Sr			0.8959										1.0571		1.1771		1.1333
( <sup>87</sup> Sr/ <sup>86</sup> Sr) i			0.71128										0.7081		0.70943		0.7099
eSr			101.3										56.21		74.9		81.8
<sup>143</sup> Nd/ <sup>144</sup> Nd	0.512247 ± 6	0.5122464 ± 7			0.5121893 ± 5				0.5122532 ± 2		0.5122408 ± 6		0.5122060 ± 8		0.512839 ± 1		0.512235 ± 45
<sup>147</sup> Sm/ <sup>144</sup> Nd	0.1239	0.1472			0.11839				0.12188		0.11906		0.11591		0.12015		0.12225
( <sup>143</sup> Nd/ <sup>144</sup> Nd) i	0.512	0.51197			0.51195				0.51201		0.51197		0.51197		0.51203		0.51199
eNd	-4.8	-5.6			-5.8				-4.6		-5.5		-5.8		-4.3		-5

Location Sample number Rock Type	Figeac CLB96-2 rhyodacite	Figeac CLB96-3 rhyodacite	Figeac CLB96-5 rhyolite	LM South CLB96-6 rhyodacite	LM South CLB96-7 rhyodacite	LM South CLB96-8 rhyodacite	LM South CLB96-9 rhyodacite	LM South CLB96-10 rhyodacite	LM South CLB96-11 rhyodacite	LM South CLB96-12 rhyodacite	LM North CLB96-13 rhyodacite	LM North CLB96-14 rhyodacite	LM North CLB96-16 rhyodacite	LM North CLB96-17 rhyodacite
SiO <sub>2</sub>	70.55	61.3	73.95	60.08	61.29	71.91	57.05	58.96	60.31	65.51	58.14	52.82	60.79	58.99
Al <sub>2</sub> O <sub>3</sub>	15.96	17.6	14.46	17.61	17.58	15.02	18.88	17.46	17.82	17.78	16.48	16.46	15.77	16.56
Fe <sub>2</sub> O <sub>3</sub>	1.79	6.31	1.11	6.6	6.22	2.16	5.71	7.21	4.48	6.93	7.09	6.24	7.55	7.17
MnO	0.01	0.05		0.04	0.06	0.03	0.06	0.01	0.03	0.1	0.07	0.05	0.06	0.02
MgO	0.61	4.02	0.33	3.43	3.46	0.57	4.1	3.82	2	3.03	6.18	6.12	7.1	6.13
CaO	0.21	5.45	0.14	0.46	0.43	0.22	1.07	0.31	0.2	0.3	6.81	0.85	0.87	0.51
Na <sub>2</sub> O	0.78	2.1	0.92	0.79	1.89	1.41	0.55	0.08	0.73	1.08	2.49	2.68	2.5	1.3
K <sub>2</sub> O	9.54	1.83	8.82	10.1	8.03	8.29	1.16	10.32	11	8.46	1.92	5.22	4.62	8.48
TiO <sub>2</sub>	0.43	0.97	0.25	0.71	0.78	0.32	0.8	0.81	0.77	0.68	0.62	0.64	0.6	0.66
P <sub>2</sub> O <sub>5</sub>	0.1	0.26	0.04	0.15	0.14	0.04	0.17	0.16	0.13	0.14	0.14	0.15	0.12	0.14
LOI	1.74	3.14	1.45	3.51	3.59	1.98	4.14	4.38	2.8	3.97	2.37	4.97	4.81	3.81
Cr	13.25	420	10.13	450	180	11.58	549	462	266.48	416	587	224	260	611
V	27.28	140	23.04	179	170	19.42	184	164	111.53	149	163	132	138	167
Sc	22.43	28	8.18	14	14	8	11	10	55.98	13	23	16	18	16
Ni	5.74	60	7.32	59	30	4.75	59	93	42.33	74	100	35	34	96
Co	4.36	19.01	1.95	20.89	20.49	2.88	28.55	8.85	14.49	18.23	29	15.89	23.37	29.42
Cu	11	14	9.86	21	23	6.36	34	224	133.59	19	22	15	11	13
Rb	238.61	32.49	207.54	192	147	172.66	171	163	201.25	201	65	127	122	173
Sr	26.02	241	12.07	44.59	80.48	34.4	32.89	26.88	34.56	42.04	228	61.94	60.17	38.7
Y	19.88	31.54	17.98	49.2	26.72	21.83	24.34	41.85	12.58	82.02	20	22.47	22.82	18.54
Zr	207.46	162	238.99	141	165	182.77	163	149	172.97	141	126	139	130	126
Nb	12.01	10.92	11.32	8.39	11.03	9.85	8.22	9.74	8.28	7.1	8.62	8.19	7	
Cs	15.92	2.98	19.73	4.8	4.41	6.22	3.83	1.61	3.1	7.89	5.83	8.3	7.54	7.84
Ba	696.92	1204	677.58	700	977	1092.59	661	1108	1436.93	1805	464	869	560	560
La	31.7	34.69	35.66	33.848	30.424	37.26	17.383	20.025	17.2	57.892	21.349	22.921	23.771	15.903
Ce	64.19	66.132	69.38	49.659	56.49	63.41	41.004	17.305	22.93	75.245	44.356	47.217	48.736	38.128
Pr	7.6	7.997	8.15	9.684	6.969	7.77	5.1341	6.096	4.19	14.809	5.325	5.475	5.616	4.264
Nd	28.03	30.806	29.61	38.391	26.419	28.33	20.71	24.723	16.4	56.588	19.953	20.545	20.76	16.19
Sm	5.22	5.947	5.43	8.272	5.188	5.34	4.485	5.214	3.31	11.477	4.077	3.942	3.87	3.255
Eu	0.61	1.318	0.73	1.767	1.246	0.79	0.844	1.455	0.81	2.158	0.988	0.871	0.832	0.757
Gd	4.44	5.419	4.29	8.216	4.534	4.52	4.467	6.286	3.14	11.978	3.715	3.454	3.481	2.972
Tb	0.6	0.826	0.66	1.27	0.705	0.69	0.661	0.843	0.48	1.794	0.578	0.543	0.553	0.473
Dy	3.83	4.715	3.58	7.396	4.111	3.92	3.721	5.197	2.7	10.134	3.449	3.329	3.323	2.766
Ho	0.78	0.987	0.74	1.612	0.842	0.8	0.752	1.117	0.52	2.004	0.72	0.715	0.712	0.574
Er	2.23	2.559	2.13	3.982	2.36	2.21	1.927	3.0059	1.4	4.893	1.198	2.038	1.952	1.532
Tm	0.32	0.37	0.33	0.59	0.36	0.33	0.28	0.49	12	0.69	0.35	0.32	0.3	0.24
Yb	2.09	2.339	2.23	3.2	2.258	2.16	1.669	2.711	1.26	3.06	1.93	1.935	1.862	1.432
Lu	0.33	0.344	0.35	0.466	0.344	0.32	0.24	0.409	0.19	0.443	0.294	0.301	0.285	0.211
Hf	6.18	4.35	7.82	3.91	4.35	6.42	4.44	3.93	5.61	3.97	3.47	3.88	3.61	3.46
Ta	1.98	0.84	1.72	0.66	0.81	2.02	0.78	0.65	1.36	0.69	0.55	0.71	0.65	0.57
Pb	63.67	208	13.96	9.91	9.53	9.24	8.5	16.48	10.515	9.69	18	12.2	10.05	11.37
Th	14.47	10.17	15.73	9.85	9.24	16	9.82	7.8	10.18	8.85	7.83	8.96	8.33	7.8
U	5.65	3.06	4.61	2.9	2.85	4.27	3.12	2.68	3.93	2.68	2.59	2.33	2.59	2.67
Age: 300 Ma														
<sup>87</sup> Sr/ <sup>86</sup> Sr		0.710526 ± 10		0.7423544 ± 17	0.7267673 ± 20		0.7449414 ± 16	0.7476765 ± 13			0.703057 ± 170.7309075 ± 13	0.732198 ± 13	0.745231 ± 10	
<sup>87</sup> Rb/ <sup>86</sup> Sr		0.39015			5.2958						0.82667	5.9456	5.88037	
( <sup>87</sup> Sr/ <sup>86</sup> Sr) i		0.70886			0.70415						0.70704	0.70552	0.70709	
eSr		66.9			0.17						325.1	19.6	41.9	
<sup>143</sup> Nd/ <sup>144</sup> Nd	0.512235 ± 4	0.512184 ± 3	0.51213564 ± 5	0.512247 ± 3	0.512207 ± 4	0.5122009 ± 5		0.512303 ± 3	0.512280 ± 5	0.512260 ± 3	0.512268 ± 5	0.512237 ± 6	0.512246 ± 5	0.512209 ± 18
<sup>147</sup> Sm/ <sup>144</sup> Nd	0.112575	0.117458	0.11086	0.130025	0.1187	0.11394		0.12744	0.12207	0.1226	0.12352	0.11598	0.11268	0.12153
( <sup>143</sup> Nd/ <sup>144</sup> Nd) i	0.512013	0.511953	0.51213	0.51199	0.51197	0.51198		0.51205	0.51204	0.51202	0.51202	0.512	0.51202	0.51197
eNd	-4.6	-5.8	-2.21	-5.1	-5.42	-5.2		-3.9	-4.1	-4.5	-4.4	-4.7	-4.4	-5.5